

IN THE CLAIMS

1-3. (cancelled)

4. (twice amended) A method for screening for testing treatments of neurodegenerative disease comprising:

inducing protein aggregation in a first sample comprising NACP/α-synuclein by exposing a first sample to an oxidizing agent of amyloidogenic proteins;

exposing the first sample amyloidogenic proteins to a treatment;

inducing protein aggregation in a second sample comprising NACP/α-synuclein by exposing a second sample to the oxidizing agent;

measuring an aggregation level of NACP/α-synuclein in the first sample and the second sample;

and measuring comparing the aggregation level of NACP/α-synuclein in the first sample and with the aggregation level of NACP/α-synuclein in the second sample to test for a decrease in aggregation, wherein the decrease less aggregation in the first sample is indicative of an effective treatment.

5. (twice amended) The method of claim 4 44, wherein the oxidizing agent comprises oxidative stress is stimulated by a mixture of metal-ions and hydrogen peroxide.

6. (twice amended) The method of claim 4 44, wherein the oxidizing agent comprises a product of oxidative stress is stimulated using an iron-catalyzed oxidative reaction.

7. (twice amended) The method of claim 4, wherein the treatment comprises expression of a non-amyloidogenic protein that modulates aggregation of NACP/α-

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synuclein.

8. (original) The method of claim 7, wherein the non-amyloidogenic protein is β -synuclein.

9-11. (cancelled)

12. (previously presented) The method of claim 4 14, wherein the treatment comprises an agent to promote the expression of anti-amyloidogenic proteins.

13. (previously presented) The method of claim 12, wherein the anti-amyloidogenic protein is β -synuclein.

14. (new) The method of claim 4, wherein the first sample comprises cells that express NACP/ α -synuclein.

15. (new) The method of claim 14, wherein the cells are neuronal cells.

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